

FORM PTO-1449(Modified)	ATTY. DOCKET NO.: V0179/7000	SERIAL NO.: 09/463874
	LIST OF PATENTS AND PUBLICATIONS FOR APPLICANT'S INFORMATION DISCLOSURE STATEMENT	
	APPLICANT: Wanker et al.	
	FILING DATE: Herewith	GROUP: Not yet assigned

## U.S. PATENT DOCUMENTS

Exam Init	Ref Des	Document No.	Date	Name	Class	Sub Class	FILING DATE If Appropriate
OC		5,234,814	08/10/93	Card et al.	435	7.21	06/01/89
OC		5,723,301	03/03/98	Burke et al.	435	7.1	11/03/95

## FOREIGN PATENT DOCUMENTS

	Country & Doc. No. (11)	Pub. Date (43)	Class	Sub Class	Translation Yes No
	EP 0 206302A2	30.12.86			
	EP 0 293 249 A1	30.11.88			
	EP 0 854 364 A1	22.07.98			
OC	WO95/29243	02.11.95			
	WO96/12544	02.05.96			
	WO96/28471	19.09.96			
	WO97/17445	15.05.97			

## OTHER ART

(Including Author, Title, Date, Pertinent Pages, Publication, Etc.)

OC		Kisilevsky et al., "Arresting amyloidosis <i>in vivo</i> using small-molecule anionic sulphonates or sulphates: Implications for Alzheimer's disease", (2/95), pp. 143-148, <i>Nature Medicine</i> , Vol. 1, No. 2.
OC		Trottler et al., "Polyglutamine expansion as a pathological epitope in Huntington's disease and four dominant Cerebellar ataxias", (11/23/85), pp. 403-406, <i>Nature</i> Vol. 378.
OC		Gutekunst et al., "Identification and localization of huntingtin in brain and human lymphoblastoid cell lines with anti-fusion protein antibodies", (9/95), pp. 8710-8714, <i>Pro. Natl. Acad. Sci. USA</i> Vol. 92 Neurobiology.
OC		M.F. Perutz, "Glutamine repeats and inherited neurodegenerative diseases: molecular aspects", (1996) pp. 848-858, <i>MRC Laboratory of Molecular Biology</i> .

EXAMINER <i>Chung</i>	DATE CONSIDERED 01/23/03
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EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant

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**OTHER ART**  
(Including Author, Title, Date, Pertinent Pages, Publication, Etc.)

		Mangiarini et al., "Exon 1 of the <i>HD</i> gene with an expanded CAG repeat is sufficient to cause a progressive neurological phenotype in transgenic mice", (11/01/96), pp. 493-506, <i>Cell</i> , Vol. 87.
		Stott et al., "Incorporation of glutamine repeats makes protein oligomerize: implications for neurodegenerative Diseases", (7/95), pp. 6509-6513, <i>Pro. Natl. Acad. Sci. USA</i> , Vol. 92, Biochemistry.
		Merlini et al., "Interaction of the anthracycline 4'-iodo-4'-deoxydoxorubicin with amyloid fibrils: Inhibition of amyloidogenesis", (3/95), pp. 2959-2963, <i>Proc. Natl. Acad. Sci. USA</i> , Vol. 92, Medical Sciences.
OC		Scherzinger, et al., "Huntingtin-encoded polyglutamine expansions form amyloid-like protein aggregates <i>in vitro</i> and <i>in vivo</i> ", (8/8/97), pp. 549-558, <i>Cell</i> , Vol. 90.
		Davies et al., "Formation of neuronal intranuclear inclusions underlies the neurological dysfunction in mice Transgenic for the HD mutation", (8/8/97), pp. 537-548, <i>Cell</i> , Vol. 90.
		Derwent, "Nucleic acid fragments associated with spinocerebellar ataxia type 2-contain increased number of CAG repeat region compared to normal gene", (5/7/98), XP-002101183.
		Tateishe et al., "Removal of causative agent of creutzfeldt-jakob disease (CJD) through membrane filtration method", (1993), pp. 357-362, <i>Membrane</i> , 18(6).

EXAMINER <i>Clampson</i>	DATE CONSIDERED <i>07/23/03</i>
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